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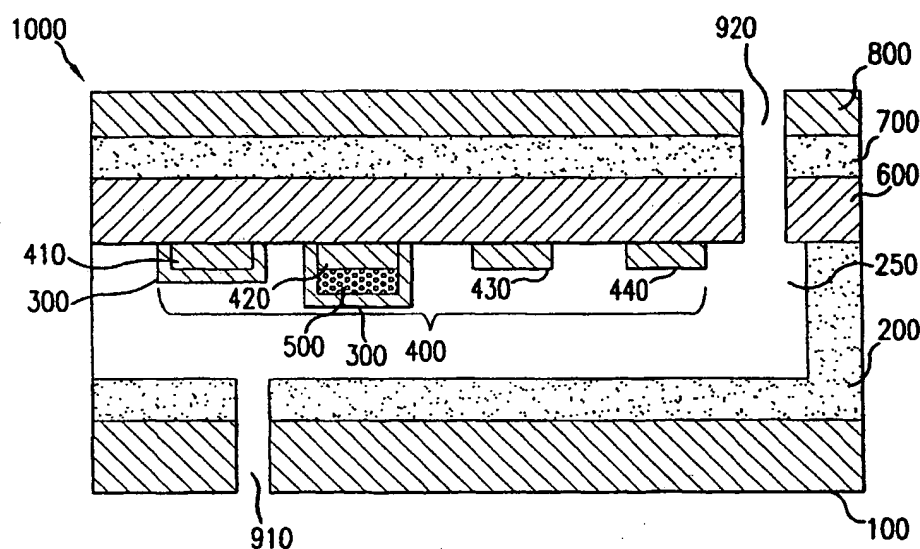
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(54) Title: ASSAY DEVICE FOR MEASURING CHARACTERISTICS OF A FLUID ON A CONTINUAL BASIS



(57) Abstract: The present invention is directed at an assay device for detecting and enabling measurement of an analyte in a fluid. The assay device contains: a) an inlet port to receive fluid; b) a well in fluid communication with the inlet port; c) an outlet port in fluid communication with the well, wherein the outlet port is designed to allow discharge of the fluid; d) at least one first working electrode and at least one reference electrode disposed within the well; e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and f) at least one membrane disposed over or around the reactant to regulate contact of the analyte in the fluid with the reactant.

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**ASSAY DEVICE FOR MEASURING CHARACTERISTICS  
OF A FLUID ON A CONTINUAL BASIS**

**CROSS REFERENCE TO RELATED APPLICATIONS**

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This application claims priority to U.S. Provisional Applications Serial No. 60/128,198 filed April 7, 1999; Serial No. 60/139,975 filed June 18, 1999; Serial No. 60/139,976 filed June 18, 1999; Serial No. 60/165,809 filed November 16, 1999; and Serial No. 60/182,698 filed February 15, 2000, all of which are

10 incorporated by reference.

**FIELD OF INVENTION**

15       The present invention relates in general to analyte detection systems and methods. More specifically, this invention relates to an assay device that detects the presence, amount or other characteristic of an analyte of interest such as a glucose level within a fluid collected from a tissue on a continuous/continual basis.

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**BACKGROUND OF THE INVENTION**

Medical studies have shown that the serious complications of diabetes can be significantly reduced by the proper control of the blood glucose levels. As a  
25 result, millions of diabetics monitor their blood glucose level on a daily basis via the traditional method of finger pricks and placing a blood sample into a testing apparatus. Some diabetics must monitor their blood glucose level more than just once a day. These individuals would greatly benefit from a system that continuously monitors blood glucose level without multiple finger pricks.

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Attempts have been made to simplify the testing process and eliminate the need for blood. One method has been to illuminate the skin of the individual to determine the glucose level. Unfortunately, these attempts have failed to produce a viable product for continuous monitoring of blood glucose levels.

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Another system under investigation is disclosed in U. S. Patent No. 5,961,451 to Reber et al. This patent sets forth a system which monitors the glucose level in the patient's interstitial fluid via an electrochemical assay device. However, this system is for one-time use only. The assay device must be replaced after every use. Similarly, U. S Patent No. 5,391,250 to Cheney II et al. and U. S. Patent No. 5,437,999 to Diebold et al. teach methods for fabricating electrochemical devices for one-time use biological applications.

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Existing electrochemical testing systems have certain drawbacks to the individual user as these systems are usually expensive and inaccurate. In addition, these systems often have difficulties detecting low levels of analyte present in the interstitial fluid. Also, many of the previous systems are far too large for the individual user to use on a regular or continuous basis throughout the day.

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Therefore, it would be advantageous to develop an analyte assay device that is useful to continuously monitor blood glucose levels.

### SUMMARY OF THE INVENTION

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The present invention relates to an assay device for detecting an analyte in a fluid, comprising: a) an inlet port to receive fluid; b) a well in fluid communication with the inlet port; c) an outlet port in fluid communication with the well to discharge fluid; d) at least one first working electrode and at least one reference electrode disposed within the well; e) a quantity of reactant that reacts

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with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and f) at least one membrane disposed over or around the reactant to regulate contact of the analyte in the fluid with the reactant. The membrane serves to extend the useful  
5 life of the assay device by slowing consumption of the reactant. As a result, the assay device is well suited for continuous monitoring applications.

In addition, the present invention relates to an assay device for detecting and enabling measurement of an analyte in a fluid comprising: a) an inlet port to  
10 receive fluid; b) a well in fluid communication with the inlet port; c) an outlet port in fluid communication with the well to discharge fluid; d) at least one first working electrode and at least one reference electrode disposed within the well; e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first  
15 working electrode; and f) calibration port that is in fluid communication with the well.

Advantages of the invention will be obvious from the description, or may be learned by practice of the invention. Additional advantages of the invention  
20 will be realized and attained by means of the elements and combinations particularly pointed out in the appended claims. It is to be understood that both the foregoing general description and the following detailed description are exemplary and explanatory of preferred embodiments of the invention, and are not restrictive of the invention, as claimed.

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The accompanying drawings, which are incorporated in and constitute a part of this specification, illustrate preferred and alternate embodiments of the invention and together with the description, serve to explain the principles of the invention.

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## BRIEF DESCRIPTION OF THE DRAWINGS

**Figure 1** is a cross-section view of a preferred embodiment of the assay device according to the present invention.

5        **Figure 2** is a cross-section view of an alternative embodiment of the assay device.

**Figure 3** is a cross-section view of an another alternative embodiment of the assay device.

10       **Figure 4** depicts an exploded view of still another embodiment of an assay device according to the present invention.

## DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

15        The present invention may be understood more readily by reference to the following figures and their previous and following description, including the detailed description of the invention and the examples provided herein. It is to be understood that this invention is not limited to the specific devices and methods  
20       described, as specific device components and/or process conditions as such may, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting.

25        It must also be noted that, as used in the specification and the appended claims, the singular forms "a," "an" and "the" comprise plural referents unless the context clearly dictates otherwise. For example, reference to a component in the singular is intended to comprise a plurality of components.

As used herein, "analyte" shall mean the component that is being detected or measured in an analysis. In particular, the analyte may be any chemical or biological material or compound suitable for passage through a biological membrane technology known in the art, of which an individual might want to know the concentration or activity inside the body. Glucose is a specific example of an analyte because it is a sugar suitable for passage through the skin, and individuals, for example those having diabetes, might want to know their blood glucose levels. Other examples of analytes include, but are not limited to, such compounds as sodium, potassium, bilirubin, urea, ammonia, calcium, lead, iron, lithium, salicylates, pharmaceutical compounds, and the like.

Ranges may be expressed herein as from "about" or "approximately" one particular value and/or to "about" or "approximately" another particular value. When such a range is expressed, another embodiment comprises from the one particular value and/or to the other particular value. Similarly, when values are expressed as approximations, by use of the antecedent "about," it will be understood that the particular value forms another embodiment.

The present invention is directed at an assay device for detecting and enabling measurement of an analyte in a fluid. The assay device contains: a) an inlet port to receive fluid; b) a well in fluid communication with the inlet port; c) an outlet port in fluid communication with the well, wherein the outlet port is designed to allow discharge of the fluid; d) at least one first working electrode and at least one reference electrode disposed within the well; e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and f) at least one membrane disposed over or around the reactant to regulate contact of the analyte in the fluid with the reactant.

Moreover, the present invention is directed to an assay device containing:

- a) an inlet port to receive fluid; b) a well in fluid communication with the inlet port; c) an outlet port in fluid communication with the well, wherein the outlet port is designed to allow discharge of the fluid; d) at least one first working  
5 electrode and at least one reference electrode disposed within the well; e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and f) calibration port that is in fluid communication with the well.

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The assay device according to the present invention is suitable for use in a continuous/continual analyte monitoring system, such as that disclosed in International Application No. PCT/US99/16378, entitled "System and Method for Continuous Analyte Monitoring," filed July 20, 1999, which is incorporated  
15 herein by reference.

Referring now to **Figure 1**, a preferred embodiment of the assay device **1000** according to the present invention is shown. In this embodiment, the assay device **1000** comprises a bottom layer **100** that is in fluid communication with a  
20 channel-forming adhesive layer **200** via an inlet port **910**. The channel-forming adhesive layer **200** has a layer of adhesive with a channel cut into it to form a well **250**. Within the well **250**, are the membrane **300** and electrodes **400**.

In the preferred embodiment shown in **Figure 1**, there are four electrodes  
25 **400**: electrode **410** is a working electrode, electrode **420** is a working electrode, electrode **430** is a reference electrode and electrode **440** is a counter-electrode. At least one of the working electrodes **420** is coated with a reactant **500**. The electrodes **400** are disposed on or in a support base **600**. The support base **600** is adjacent to an adhesive layer **700**. An outlet port **920** extends through the support  
30 base layer **600**, the adhesive layer **700**, and a top layer **800**.



The bottom layer 100 provides structural support to the assay device 1000 and serves as the interface between the fluid source and the assay device 1000. Any suitable material, in any thickness or shape may be used for the bottom layer  
5 100. Example suitable materials include acrylic, polyester, plastic, ceramic, polycarbonate and polyvinylchloride.

The inlet port 910, which provides fluid communication between the bottom layer 100, the channel-forming adhesive layer 200, and the well 250, may  
10 be in any position and in any dimension/shape to allow sufficient flow to the electrodes 400. The inlet port 910 is suitable for alignment with holes/porations in a tissue from which fluid is to be drawn, such as interstitial fluid. An example of a mechanism to facilitate alignment of the assay device 1000 with the holes/porations in the tissue is disclosed in U.S. Provisional Application Serial  
15 No. 60/140,257 filed June 18, 1999 entitled "System and Method for Alignment of Micropores for Efficient Fluid Extraction and Substance Delivery," which is incorporated herein by reference.

The channel-forming adhesive layer 200 forms the well 250 to limit the  
20 volume of fluid within the assay device 1000. Suitable materials for the channel-forming adhesive layer 200 are compatible with the fluid of interest, provide adhesive support to the assay device 1000, and are thick enough to provide a well 250 from a channel cut into the channel-forming adhesive layer 200. Preferably, the fluid of interest is blood or interstitial fluid, thereby requiring the channel-  
25 forming adhesive layer 200 to be constructed from adhesive-like materials that are not water-soluble.

The electrodes 400 are disposed on or in a support base 600 using screen-printing, pad printing, sputter coating, photolithography or other suitable  
30 techniques, using known inks and dielectrics. The support base 600 may be of

any thickness effective to provide support and bind the electrodes 400. One preferred embodiment includes a support base 600 of 10 mil thick transparent polyester. Other suitable materials may be used including ceramic, polycarbonate, and polyvinylchloride.

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Moreover, an adhesive layer 700 and a top layer 800 provide additional support to the assay device 1000. The adhesive layer 700 binds the support base 600 to the top layer 800. The material of construction and dimension of the adhesive layer 700 is not critical to the present invention, thereby allowing any effective adhesive to be used. The top layer 800, like the bottom layer 100, provides structural support to the assay device 1000. Preferably, the top layer 800 is constructed of the same material or compatible material as the bottom layer 100.

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An outlet port 920 allows discharge of the fluid from the well 250 through the support base 600, adhesive layer 700, and the top layer 800. It may be in any position and in any dimension/shape to allow sufficient flow to the electrodes 400. The outlet port 920 also is suitable for connection to a supply of vacuum sufficient to draw fluid through the well 250. In one preferred embodiment, the vacuum is sufficient to produce fluid from the skin at a site of where small holes/porations have been made in the tissue. The well 250 serves to expose the membrane 300 and electrodes 400 to the fluid that is monitored. Therefore, the well 250 is preferably of a dimension that the membrane 300 and the electrodes 400 do not obstruct the flow of fluid.

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As explained in one example hereinafter, a reactant 500 reacts with the analyte to form a reaction product. The reaction product is in fluid communication with one or both of the working electrodes 410 and 420 whereby electrons are created. Depending on the reactant's composition, the reactant 500 may react with glucose, which may in turn form hydrogen peroxide. In this

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embodiment, when hydrogen peroxide contacts a working electrode, oxygen gas, hydrogen ions and electrons are produced.

Each working electrode 410 and 420 may be made from a variety of materials such as carbon and metals such as gold or silver. Preferably, each working electrode 410 and 420 is made from catalytic metals such as platinum, palladium, chromium, ruthenium, rhenium, or mixtures thereof. Most preferably, the working electrodes 410 and 420 contain platinum.

To detect and/or measure the level of an analyte present in a fluid, at least one working electrode and at least one reference electrode are necessary. However, more than one working electrode and one or more counter-electrodes may also be present. For example, in the embodiment shown in Figure 1, the working electrode 410 does not contain the reactant and therefore it produces an electrical signal that is indicative of the fluid without the analyte. This allows reduction or elimination of the signal due to various interferent compounds by subtracting the electrical signal of the working electrode 410 from the electrical signal of the working electrode 420.

Alternatively, one working electrode may be used if the levels of interference are not significant or if an interference blocking layer is included. This interference blocking layer could be positioned anywhere between the fluid to be analyzed and the working electrodes 410 and 420. In one preferred embodiment, the interference blocking layer is placed directly over the working electrodes 410 and 420. In another preferred embodiment, the interference blocking layer is placed adjacent to the membrane 300. Suitable interference blocking layers include NAFION™ and cellulose acetate. Possible interferents include: acetaminophen, ascorbic acid, unconjugated bilirubin-, cholesterol, creatinine, dopamine, gentisic acid, heparin, ibuprofen, salicylate, tetracycline, tolbutamide, triglycerides and uric acid.

The reference electrode 430 establishes a potential relative to the fluid. Preferably, the reference electrode 430 contains silver/silver-chloride. The counter-electrode 440, which is optional, serves to ground the current generated by the working electrodes 410 and 420. Preferably, the counter-electrode 440 contains substantially the same materials as the working electrodes 410 and 420. The assay device 1000 may contain more than one working electrode, more than one reference electrode and more than one counter-electrode, as is well known in the art.

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The active surface of the electrodes 400 may be any shape and dimension to effectively operate. Particularly, the surface area of the any of the electrodes 400 can be varied as long as there is sufficient sensitivity to measuring the current. Preferably, the electrodes 400 have active surface areas between 0.1 mm<sup>2</sup> and 10 mm<sup>2</sup>. Most preferably, the electrodes 400 have a surface area of 1mm<sup>2</sup>.

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After the assay device 1000 is constructed, the working electrodes 410 and 420 may be pre-conditioned by running at a specific voltage, such as +1.6 V relative to the reference electrode 430 for a suitable amount of time, such as 30 minutes, in a buffer system. This conditions the surface of the working electrodes 410 and 420 and increases their sensitivity to the reaction product generated by the reactant 500. Alternately, the working electrodes 410 and 420 could be conditioned for shorter times at higher voltages.

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The working potential will depend on the composition and shape of catalytic surface area. As such, the working potential can vary from 200 mV to 2 V. Such potential may be supplied via a monitoring unit coupled to the assay device wherein the monitoring unit utilizes an amperometric or coulometric measurement technique, known in the art. The working potential is generated

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either by holding the working electrodes 410 and 420 at a positive potential or by holding the counter-electrode 440 at a negative potential. For example, the working electrodes 410 and 420 may be held at +800 mV and the reference electrode 430 and counter-electrode 440 at 0 mV, or the working electrodes 410 and 420 may be held at 0 mV and the reference electrode 430 and counter-electrode 440 at -800 mV.

The electrodes may be connected to leads that in turn are connected to a monitoring unit (Figure 4), patient worn or otherwise, via traces of graphite or silver/silver-chloride. However, other conductive material such as gold or tin are suitable to connect the electrodes to the leads. These traces could be applied via any method that provides a sufficient resolution such as ink-jet printing or pad printing. In addition, the printed traces could be replaced with traditional connection techniques.

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A quantity of reactant 500 that reacts with the analyte to form a reaction product is disposed proximate to the at least one first working electrode such that when the analyte contacts the reactant 500, the reaction product is in fluid communication with the working electrode. Preferably, the quantity of reactant 500 covers a portion of the first working electrode (working electrode 420 shown in Figure 1). The quantity of reactant 500 may also be disposed on or in at least one working electrode. The reactant 500 is selected to react with a specific analyte. In one preferred embodiment, the quantity of reactant 500 is suitable to react with glucose. As such, suitable reactants for the analyte glucose include glucose oxidase enzyme ("GOX"), glucose dehydrogenase ("GDH"), or mixtures thereof.

When the reactant 500 is chosen from this group, the glucose in the fluid makes contact with the reactant(s) to produce reaction products, which in the case of GOX, are gluconolactone and hydrogen peroxide. The hydrogen peroxide

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diffuses to the working electrode 420 and reacts with the catalytic metal to produce electrons as described above. Alternatively, the reactants may include a mediator as an electron receptor instead of using oxygen. In such an embodiment, the mediator reacts with the working electrode 420 to produce  
5 electrons. Mediators that are commonly used are ferrocene, ferrocyanide and their derivatives.

In one preferred embodiment, the reactant 500 is prepared by mixing 8 mg/mL GOX with 60 mg/mL bovine serum albumen ("BSA") that is dissolved in  
10 phosphate buffered saline ("PBS") that contains 10% glycerol and 0.01%  $\text{NaN}_3$ . In this preferred embodiment, 20  $\mu\text{L}$  of 25% glutaraldehyde is added to the mixture immediately before application. Preferably, a 1  $\mu\text{L}$  drop of this mixture is placed onto one of the working electrodes. The mixture is then allowed to solidify and cure at room temperature for approximately eight to sixteen hours.

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The BSA serves as a carrier for the GOX due to its multiple cross-linking sites. As such, it can be replaced with any material that has multiple surface amine groups. In addition, the combination of BSA and glutaraldehyde as a cross linking system can be replaced with a system that will immobilize the active  
20 enzyme (in this case, GOX or GDH) without inhibiting its activity. Suitable replacements include other cross-linkers, polymer films, avidin-biotin linkages, antibody linkages, and covalent attachment to colloidal gold or agarose beads.

In this preferred embodiment, the PBS acts to maintain the reactant in a  
25 neutral pH range (such as a pH of about 6.5 to about 7.5). Any suitable buffer may be used. Example buffers include phosphate, citrate, Tris-HCl, MOPS, HEPES, MES, Bis-Tris, BES, ADA, ACES, MDPSO, Bis-Tris Propane, and TES. The glycerol serves to prevent the reactant 500 from becoming dehydrated which reduces the wetting time for later use. To this end, any suitable additive

may be used. The  $\text{NaN}_3$  acts as an anti-bacterial agent. The  $\text{NaN}_3$  could be replaced by any anti-microbial agent including antibiotics and detergents.

The glutaraldehyde is a cross-linking agent that links the GOX and BSA  
5 into a matrix that will not dissolve or move from the electrode surface. In one preferred embodiment, the glycerol could be present in an amount of 5% to 50% (by weight). Alternatively, the glycerol can be replaced or supplemented by any hygroscopic preservative or wetting agents including mild detergents such as TWEEN-20™, SPAN™, TRITON™, BRIJ™, MYRJ™ and PLURONICS™  
10 families of detergents.

The proportions of any of the components to the reactant 500 and the overall amount of reactant 500 are not critical to the present invention as long as the amount is effective. These proportions and overall amount of reactant 500  
15 are limited by the preference of having an excess of reactant 500 as well as maintaining the reactant's solubility in the available volume. Preferably, the concentration of the reactant 500 is a minimum of 1 mg/mL.

The reactant 500 may be applied to a working electrode with any method  
20 that allows for volume and position control capable with techniques such as screen printing, ink-jet printing, air brush, and pad printing. For example, for application methods that use a nozzle or a screen that must continuously pass solution, the reactant 500 is preferably applied without glutaraldehyde, and then, the glutaraldehyde is placed down. This avoids fouling the nozzle with  
25 solidifying material. Once applied, the reactant 500 is dried and cured with the times of each varying based upon the amount and thickness of reactant layer(s) and the composition. Suitable drying conditions include temperatures up to 150°C, controlled humidity, and cure times of 15 minutes to 24 hours.

The GOX enzyme will saturate at concentrations of approximately 3 mM glucose. In order to detect higher levels of glucose, the concentration reaching the reactant 500 must be held to a fraction of the total concentration. To accomplish this, a membrane 300 is disposed over or around the reactant 500. In one preferred embodiment, the membrane 300 is disposed over or around all of the electrodes 400 as shown in Figure 3. In another preferred embodiment, the membrane 300 is disposed over or around the working electrodes 410 and 420 as shown in Figure 2. Alternatively, the membrane 300 is disposed over or around each electrode 400 or each working electrode 410 and 420 as shown in Figures 1.

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The membrane 300 preferably is a diffusion-limiting membrane that extends the linear range and the lifetime of the assay device 1000 system and makes it useful in a continuous/continual monitoring system. The membrane 300 has pores which regulate diffusion of an analyte therethrough. Therefore, the membrane 300 may be sized to limit the rate at which the analyte or an interferent makes contact with the reactant 500, thereby increasing the linear range of the assay device 1000. For example, the membrane 300 may have low porosity to reduce glucose flux. As such, the membrane 300 limits the amount of analyte that is present at the electrodes at any one time, allowing the electrodes 400 to operate continuously over long periods of time without depleting the reactant. In fact, a monitoring unit coupled to the at least one working electrode may continuously draw fluid through the assay device and detect the presence or level of an analyte in excess of 24 hours, more preferably in excess of 48 hours, and still more preferably in excess of 70 hours.

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One preferred membrane 300 is a 0.01  $\mu\text{m}$  pore diameter polycarbonate ("PC") track-etch member 6  $\mu\text{m}$  thick. Other suitable membranes that effectively produce a diffusion rate include dialysis membranes, polyurethane membranes, or polyvinylchloride membranes. Castable membranes such as NAFION™,



cellulose acetate, silastics and alkoxy silanes are also effective for this use.

Additionally, multiple membranes may be used.

5       The membrane 300 may be secured by a layer of cross-linked BSA in the same buffer as the reactant 500. In one preferred embodiment, the cross-linked BSA consists of 60 mg/mL BSA dissolved in PBS containing 10% glycerol and 0.01% NaN<sub>3</sub>. In addition, 20 µL/mL of 25% glutaraldehyde may be added immediately before use.

10       Any effective amount of this layer of cross-linked BSA may be used. In one preferred embodiment, a 2 µL drop of the cross-linked BSA is placed on the reactant covered electrode. Then, a 5 mm diameter circle of the 0.01 µm PC membrane is placed over the drop of cross-linked BSA or other suitable, large polyamine. The membrane 300 is gently pressed into place under a sheet of  
15       parafilm. It is allowed to cure for 16 hours at room temperature under the parafilm, and then the parafilm is removed.

      The membrane 300 may be disposed over or around the reactant 500 by any suitable method including lamination, gluing, pressing, rolling and stretching.  
20       Any such method should not be destroyed by the fluid to be collected and analyzed. However, if an additional component is added, such as glue or other adhesion material, the additional component should be permeable in the fluid, such as Nafion is permeable in an aqueous-based fluid. Other suitable adhesives include epoxies, UV curable adhesives, pressure sensitive adhesives and  
25       hydrogels such as HEMA.

      In operation, the assay device 1000 is positioned on a tissue site overlying one or more openings made in the tissue. The openings in the tissue may be made by a variety of means, such as those disclosed in commonly assigned U.S.  
30       Patent No. 5,885,211. Fluid enters the assay device 1000 of **Figure 1** through the

inlet port 910. Under application of vacuum at the outlet port 920, the fluid travels through the well 250 and contacts the membrane 300. The membrane 300 is permeable to the analyte in the fluid and thereby allows the analyte to contact the electrodes 400 and the reactant 500. The analyte reacts with the reactant 500 to generate a reaction product. The reaction product contacts the working electrode 420 and creates electrons, thereby generating a current flow. The fluid continues through the outlet port 920 whereupon it exits the assay device 1000. Current flow across the working electrode(s) is measured, and from this a measurement of the analyte is obtained. The assay device 1000 may be used in conjunction with amperometric and coulometric measuring techniques.

In an amperometric measurement the current (charge/second) is measured at the applied voltage. This can be measured continuously, which is a preferred method in a flowing system. With a coulometric technique, the total charge accumulated over a period of time is measured after a voltage is applied. Typically, the fluid is allowed to react with the reactant over a fixed period of time, thereby generating a reaction product. Then a voltage is applied and the current, which is measured over a fixed period of time, is integrated (added) to calculate the total amount of charge produced by the reaction product. This alternative method has the advantage of generating larger signals and reducing the impact of electroactive interfering substances.

The membrane 300 preserves the life of the reactant 500 by helping to hold the reactant 500 in place and to thereby reduce the risk of rapid dissolution of the reactant 500 in the fluid. Also, by restricting the amount of analyte and interferents to the reactant 500, the membrane 300 helps ensure that the reactant 500 is in excess than what is needed to fully engage the analyte. In this way, as the reactant 500 degrades over time, it may remain in excess and deterioration in performance will be minimized.

Referring now to **Figure 2**, a variation of the assay device 1000 is shown in which a well 250 opens into an outlet port 920. In such a configuration, the outlet port 920 does not provide fluid communication through the support base 600, the adhesive layer 700, and the top layer 800. However, the outlet port 920  
5 is suitable for connection to a supply of vacuum sufficient to draw fluid through the well 250. The outlet port 920 may be filled with wiring and drain tubing, and then epoxy sealed.

**Figure 3** shows another alternative preferred embodiment of the assay  
10 device 1000. As in **Figure 2**, the well 250 is coupled to the outlet port 920. However, **Figure 3** shows an assay device that comprises only a support base 600 and bottom layer 100. The bottom layer 100 provides the well 250 to expose the membrane 300 and electrodes 400 to the fluid.

**Figure 4** shows another embodiment of an assay device 1000 according  
15 to the present invention. This embodiment includes a bottom layer 100, a channel-forming adhesive layer 200, a support base 600, an adhesive layer 700, and a top layer 800. **Figure 4** also includes an inlet port 910, a well 250, an outlet port 920, a calibration port 950, and drain tubing 940. Although not  
20 shown, this embodiment includes at least one working electrode, a reference electrode, and a reactant proximate to the working electrode as shown in **Figures 1-3**. This embodiment may optionally include at least one membrane, as shown in **Figure 1**. The electrodes are connected to a monitoring unit 970 via leads 960. The monitoring unit 970 is also connected to assay device 1000 via the drain  
25 tubing 940. The drain tubing 940 provides vacuum to the assay device 1000.

Similar to the assay device 1000 of **Figure 2**, the well 250 shown in **Figure 4** opens into the outlet port 920. The calibration port 950 is suitable for connection to a reservoir 980 containing calibration fluid. In addition, the

calibration port 950 may include a membrane 990 permeable to the calibration fluid.

In one embodiment, the calibration fluid consists of water and the analyte  
5 to be detected. Other compounds may also be present, such as surfactants, which  
ensure smoother flow by reducing surface tension, such as SDS, or any of the  
detergents herein described. Additionally, preservatives such as azide, EDTA, or  
any antibacterial or appropriate biocide that will not degrade or interfere with the  
reactant's performance may be added to the calibration fluid. Moreover, the  
10 calibration fluid may include thickeners, such as polymers and proteins, to  
simulate the flow characteristics of the analyte-containing fluid that is being  
measured.

The reservoir 980 is in fluid communication with the well 250 such that  
15 the calibration fluid flushes the well 250 to contact the electrodes with the  
calibration fluid. The calibration fluid is removed from the well 250 through the  
outlet port 920 under application of vacuum.

The reservoir 980 may release the calibration fluid into the well 250 using  
20 any effective mechanism. In one embodiment, the reservoir 980 comprises a  
bag-like member that opens and releases the calibration fluid into the well 250 in  
response to application of vacuum applied at the outlet port 920. Moreover, the  
reservoir 980 may be formed of a material that when mechanically punctured  
releases the calibration fluid into the well 250.

25

Alternatively, the membrane 990 is a self-sealing membrane that is  
rupturable to allow introduction of the calibration fluid such as by a syringe  
containing calibration fluid to deliver the calibration fluid into the well 250.  
Preferably, the calibration fluid is delivered into the well 250 while vacuum is  
30 applied at the outlet port 920. In another preferred embodiment, the calibration

fluid is introduced into the well 250 via a valve that operates as a one-way valve or is controlled external to the assay device 1000.

Throughout this application, various publications are referenced. The  
5 disclosures of these publications in their entireties are hereby incorporated by reference into this application in order to more fully describe the state of the art to which this invention pertains.

It will be apparent to those skilled in the art that various modifications and  
10 variations can be made in the present invention without departing from the scope or spirit of the invention. Other embodiments of the invention will be apparent to those skilled in the art from consideration of the specification and practice of the invention disclosed herein. It is intended that the specification and examples be considered as exemplary only, with a true scope and spirit of the invention being  
15 indicated by the following claims.

**What is claimed is:**

1. An assay device for detecting an analyte in a fluid, comprising:
  - a) an inlet port to receive fluid;
  - b) a well in fluid communication with the inlet port;
  - c) an outlet port in fluid communication with the well to discharge fluid;
  - d) at least one first working electrode and at least one reference electrode disposed within the well;
  - e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and
  - f) at least one membrane disposed over or around the reactant to regulate contact of the analyte in the fluid with the reactant.
2. The assay device of claim 1, wherein the inlet port is suitable for alignment with holes/porations in tissue from which fluid is drawn.
3. The assay device of claim 1, wherein the outlet port is suitable for connection to a supply of vacuum sufficient to draw fluid through the well.
4. The assay device of claim 1, wherein the at least one first working electrode is comprised of a catalytic metal.
5. The assay device of claim 1, wherein the at least one first working electrode is comprised of platinum, palladium, chromium, ruthenium, rubidium, or mixtures thereof.
6. The assay device of claim 1, wherein the at least one reference electrode is comprised of silver/silver-chloride.

7. The assay device of claim 1, further comprising at least one counter-electrode disposed within the well.
8. The assay device of claim 1, further comprising a second working electrode disposed within the well.
9. The assay device of claim 1, further comprising at least one counter-electrode and a second working electrode disposed within the well.
10. A monitoring system comprising the assay device of claim 1, and further comprising a monitoring unit coupled to the assay device.
11. A monitoring system comprising the assay device of claim 1, and further comprising a monitoring unit coupled to the assay device, wherein the monitoring unit generates an analyte measurement from the assay device utilizing a coulometric or amperometric measurement technique.
12. The assay device of claim 1, wherein the quantity of reactant is disposed in or on the at least one working electrode.
13. The assay device of claim 1, wherein the quantity of reactant is comprised of glucose oxidase enzyme, glucose dehydrogenase, or mixtures thereof.
14. The assay device of claim 1, wherein the quantity of reactant is suitable to react with glucose.
15. The assay device of claim 1, wherein the at least one membrane comprises pores which are sized to limit the rate at which the analyte makes contact with the reactant.

16. The assay device of claim 1, wherein the at least one membrane comprises pores which are sized to limit the rate at which an interferent makes contact with the reactant.
17. The assay device of claim 1, wherein the at least one membrane is disposed over or around the at least one first working electrode.
18. The assay device of claim 1, wherein the at least one membrane is disposed over or around the at least one first working electrode and the at least one reference electrode.
19. The assay device of claim 1, further comprising a second working electrode disposed within the well and wherein the at least one membrane is disposed over or around the at least one first working electrode and the second working electrode.
20. The assay device of claim 1, further comprising a reservoir containing a calibration fluid, wherein the reservoir is in fluid communication with the well such that the calibration fluid flushes the well and is removed from the well through the outlet port.
21. The assay device of claim 20, wherein the reservoir comprises a bag that opens and releases the calibration fluid into the well in response to application of vacuum thereto applied at the outlet port.
22. The assay device of claim 20, wherein the reservoir is formed of a material that when mechanically punctured releases the calibration fluid into the well.



23. The assay device of claim 20, further comprising a calibration port that couples the reservoir to the well, wherein the calibration port comprises a membrane permeable to the calibration fluid.
24. An assay device for detecting and enabling measurement of an analyte in a fluid, comprising:
- a) an inlet port to receive fluid,
  - b) a well in fluid communication with the inlet port,
  - c) an outlet port in fluid communication with the well to discharge fluid,
  - d) at least one first working electrode and at least one reference electrode disposed within the well,
  - e) a quantity of reactant that reacts with the analyte to form a reaction product, wherein the reaction product is in fluid communication with the at least one first working electrode; and
  - f) a calibration port that is in fluid communication with the well.

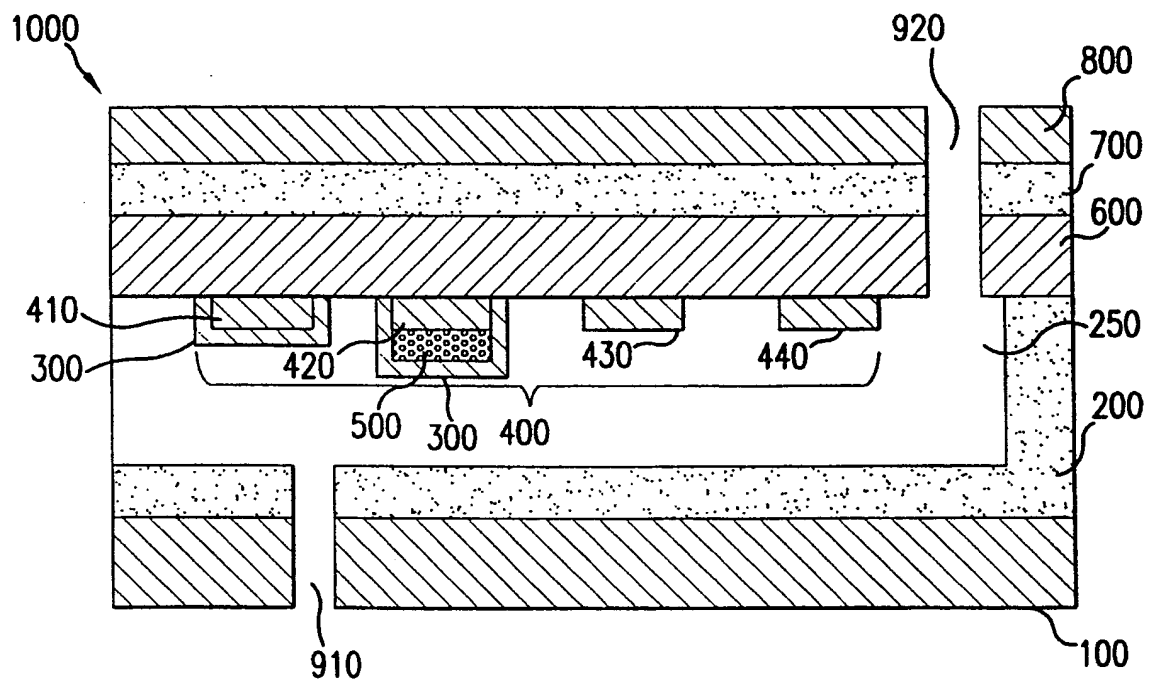


FIG. 1

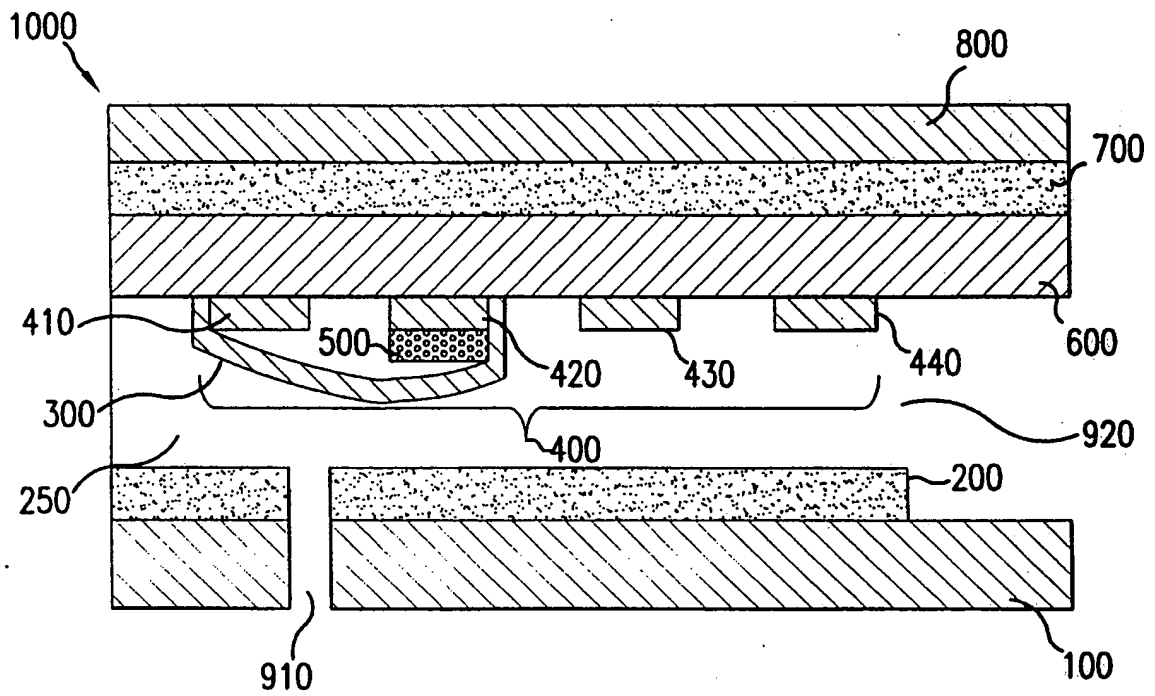


FIG.2

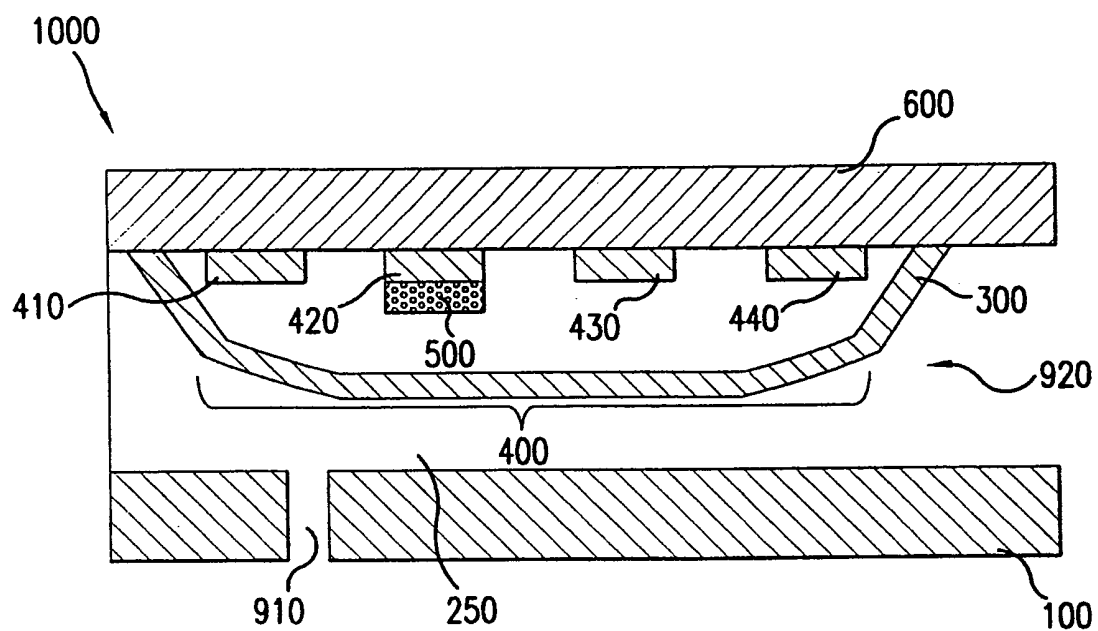


FIG.3

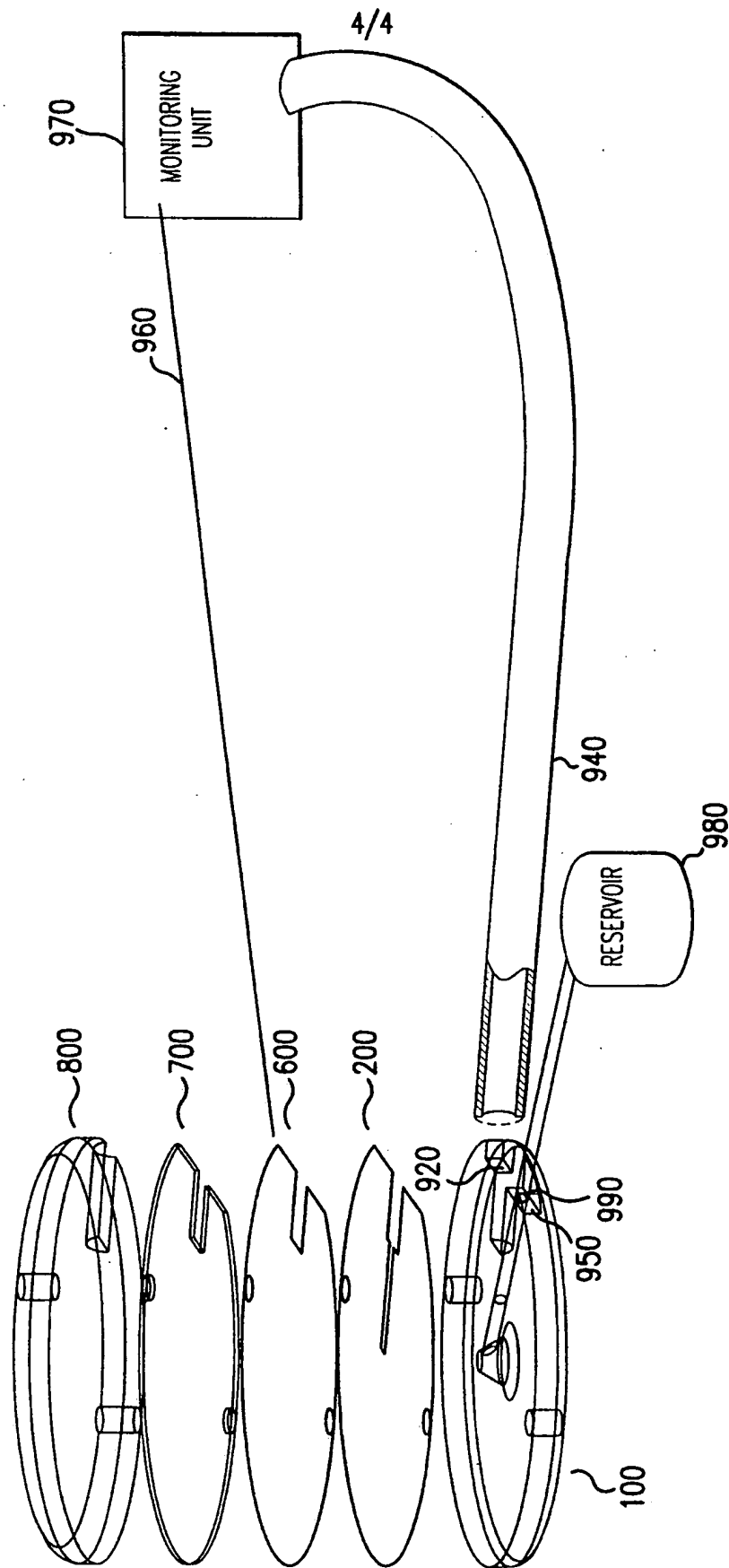


FIG. 4

# INTERNATIONAL SEARCH REPORT

International Application No.  
PCT/US 00/09393

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61B5/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 A61B 601N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)  
EPO-Internal, WPI Data

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STEINKUHL R ET AL: "MICRODIALYSIS SYSTEM FOR CONTINUOUS GLUCOSE MONITORING" SENSORS AND ACTUATORS B,CH,ELSEVIER SEQUOIA S.A., LAUSANNE, vol. B33, no. 1/03, 1 July 1996 (1996-07-01), pages 19-24, XP000632919 ISSN: 0925-4005	1,4-6, 10-14,17
A	page 20, left-hand column, line 21 -page 21, left-hand column, line 43; table 1 -/-	2,15,16, 24

☒ Further documents are listed in the continuation of box C.

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Date of the actual completion of the international search

17 July 2000

Date of mailing of the international search report

27/07/2000

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# INTERNATIONAL SEARCH REPORT

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## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	HINTSCHE R ET AL: "CHEMICAL MICROSENSOR SYSTEMS FOR MEDICAL APPLICATIONS IN CATHETERS" SENSORS AND ACTUATORS 8, CH, ELSEVIER SEQUOIA S.A., LAUSANNE, vol. B27, no. 1/03, PART 02, 1 June 1995 (1995-06-01), pages 471-473, XP000516364 ISSN: 0925-4005	1, 3-6, 11-14, 17
A	page 471, right-hand column, line 1 -page 473, left-hand column, line 2; tables 1, 2	15, 16, 24
Y	EP 0 453 283 A (TEKNEKRON SENSOR DEV CORP) 23 October 1991 (1991-10-23) column 2, line 18 -column 4, line 22; tables 1-3	1-7, 10-18, 24
Y	WO 98 30891 A (IMPLANTED BIOSYSTEMS INC) 16 July 1998 (1998-07-16)	1-7, 10-18, 24
A	page 4, line 27 -page 12, line 2; tables 2-4	20-23

# INTERNATIONAL SEARCH REPORT

Information on patent family members

Int. Patent Application No

PCT/US 00/09393

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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WO 9830891 A	16-07-1998	US 5914026 A AU 5904598 A	22-06-1999 03-08-1998



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